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Abstract: The majority of hepatocellular carcinoma (HCC) develops in the background of chronic liver inflammation caused by viral hepatitis and alcoholic or non-alcoholic steatohepatitis. However, the impact of different types of chronic inflammatory microenvironments on the phenotypes of tumors generated by distinct oncogenes is largely unresolved. To address this issue, we generated murine liver tumors by constitutively active AKT-1 (AKT) and -catenin (CAT) followed by induction of chronic liver inflammation by 3,5-diethoxycarbonyl-1,4-dihydrocollidine (DDC) and carbon tetrachloride (CCl₄). Also, the impact of DDC-induced chronic liver inflammation was compared between two liver tumor models using a combination of AKT-CAT or AKT-NRAS(G12V). Treatment with DDC and CCl₄ significantly facilitated the adenoma-to-carcinoma conversion and accelerated the growth of AKT-CAT tumors. Furthermore, DDC treatment altered the morphology of AKT-CAT tumors and caused loss of lipid droplets. Transcriptome analysis of AKT-CAT tumors revealed that cellular growth and proliferation was mainly affected by chronic inflammation and caused upregulation of Cxcl16, Galectin-3 and Nedd9 among others. Integration with transcriptome profiles from human HCCs further demonstrated that AKT-CAT tumors generated in the context of chronic liver inflammation showed enrichment of poor prognosis gene sets or decrease of good prognosis gene sets. In contrast, DDC had a more subtle effect on AKT-NRAS(G12V) tumors and primarily enhanced already existent tumor characteristics as supported by transcriptome analysis. However, it also reduced lipid droplets in AKT-NRAS(G12V) tumors. **CONCLUSION** Our study suggests that liver tumor phenotype is defined by a combination of driving oncogenes but also the nature of chronic liver inflammation. This article is protected by copyright. All rights reserved.

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Oncogenic Driver Genes and the Inflammatory Microenvironment Dictate Liver Tumor Phenotype

Matthias S. Matter,^{1,2} Jens U. Marquardt,^{1,3} Jesper B. Andersen,^{1,4} Cristina Quintavalle,² Nikolay Korokhov,¹ Jim K. Stauffer,⁵ Kosuke Kaji,¹ Thomas Decaens,¹ Luca Quagliata,² Fathi Elloumi,⁶ Tanya Hoang,² Alfredo Molinolo,⁷ Elizabeth A. Conner,¹ Achim Weber,⁸ Mathias Heikenwalder,^{9,10} Valentina M. Factor,¹ and Snorri S. Thorgeirsson¹

The majority of hepatocellular carcinoma develops in the background of chronic liver inflammation caused by viral hepatitis and alcoholic or nonalcoholic steatohepatitis. However, the impact of different types of chronic inflammatory microenvironments on the phenotypes of tumors generated by distinct oncogenes is largely unresolved. To address this issue, we generated murine liver tumors by constitutively active AKT-1 (AKT) and β -catenin (CAT), followed by induction of chronic liver inflammation by 3,5-diethoxycarbonyl-1,4-dihydrocollidine (DDC) and carbon tetrachloride. Also, the impact of DDC-induced chronic liver inflammation was compared between two liver tumor models using a combination of AKT-CAT or AKT-NRAS^{G12V}. Treatment with DDC and carbon tetrachloride significantly facilitated the adenoma-to-carcinoma conversion and accelerated the growth of AKT-CAT tumors. Furthermore, DDC treatment altered the morphology of AKT-CAT tumors and caused loss of lipid droplets. Transcriptome analysis of AKT-CAT tumors revealed that cellular growth and proliferation were mainly affected by chronic inflammation and caused up-regulation of Cxcl16, Galectin-3, and Nedd9, among others. Integration with transcriptome profiles from human hepatocellular carcinomas further demonstrated that AKT-CAT tumors generated in the context of chronic liver inflammation showed enrichment of poor prognosis gene sets or decrease of good prognosis gene sets. In contrast, DDC had a more subtle effect on AKT-NRAS^{G12V} tumors and primarily enhanced already existent tumor characteristics as supported by transcriptome analysis. However, it also reduced lipid droplets in AKT-NRAS^{G12V} tumors. **Conclusion:** Our study suggests that liver tumor phenotype is defined by a combination of driving oncogenes but also the nature of chronic liver inflammation. (HEPATOLOGY 2016; 00:000–000)

Research on hepatocellular carcinoma (HCC) has focused predominantly on the understanding of genetic and epigenetic alterations responsible for driving the neoplastic process in the liver. In recent years, key alterations in HCC tumors have been identified and used to predict patient survival and to pro-

vide possible therapeutic targets.⁽¹⁾ However, accumulating evidence suggests that tumor phenotype is not only dictated by genetic and epigenetic alterations in the tumor cells *per se* but also influenced by molecular cross-talk between tumor cells and the surrounding microenvironment.⁽²⁾ This concept is particularly relevant for

Abbreviations: AKT, AKT-1; CAT, β -catenin; CCC, cholangiocarcinoma; CCl₄, carbon tetrachloride; DDC, 3,5-diethoxycarbonyl-1,4-dihydrocollidine; GFP, green fluorescent protein; GSEA, gene set enrichment analysis; HCA, hepatocellular adenoma; HCC, hepatocellular carcinoma; IPA, Ingenuity Pathway Analysis.

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liver cancer, where the majority of HCCs (up to 90%) develop in the background of chronic liver inflammation caused by hepatitis B or C virus, alcoholic steatohepatitis, or nonalcoholic steatohepatitis.⁽³⁾ Although the recognition that the hepatic microenvironment contributes to tumor phenotype is generally accepted, the extent to which it affects HCC phenotype, including biology, remains unresolved. A direct comparison between HCC developing in noncirrhotic liver and cirrhotic liver could give some insight into the role of chronic inflammation. However, the results are difficult to interpret because compared tumors would need to have the same molecular features, such as identical mutations, which is an almost impossible requirement to meet in a clinical setting.⁽⁴⁾

Therefore, we genetically engineered liver tumors using different combinations of oncogenes with two inflammation-promoting regimens. This approach allowed us to address the impact of (1) the different microenvironment on the same tumor type and (2) the same microenvironment on different tumor types. To initiate liver tumors, we employed oncogenes known to be frequently activated in HCC: β -catenin (CAT), NRAS^{G12V}, and AKT-1 (AKT).⁽¹⁾ Chronic liver inflammation was induced by feeding mice 3,5-diethoxycarbonyl-1,4-dihydrocollidine (DDC), a nongenotoxic hepatotoxin causing portal inflammation and fibrosis.⁽⁵⁾ As a second inflammatory model, we used chronic administration of carbon tetrachloride (CCl₄), resulting in centrilobular liver damage with accompanying inflammation and fibrosis.⁽⁶⁾

Our results provide evidence that chronic inflammation driven by the DDC diet or exposure to CCl₄ had

a profound impact on liver tumor phenotype. Both inflammation-promoting regimens increased tumor growth, accelerated malignant conversion, and caused inflammation-specific changes in the tumor transcriptome of AKT-CAT tumors. DDC-mediated inflammation also influenced the biology of AKT-NRAS^{G12V}-driven hepatocarcinogenesis, albeit to a lesser degree compared to AKT-CAT tumors. We conclude that the outcome of liver cancer development is dependent on the cooperation between initiating oncogenes and the nature of inflammation.

Materials and Methods

HYDRODYNAMIC INJECTION, VECTOR CONSTRUCTS

For tumor generation, oncogenic plasmids and sleeping beauty transposase (ratio of 10:1) were diluted in 2 mL saline (0.9% NaCl) and injected into the tail vein of FVB/n female mice 6–8 weeks old (National Cancer Institute, Frederick, MD) in 5–7 seconds.⁽⁷⁾ For AKT-CAT tumors, plasmids encoding for activated AKT (myristoylated AKT) and CAT (truncated Δ 90N-CAT) were used.⁽⁸⁾ For AKT-NRAS^{G12V} tumors, AKT1 and N-Ras^{G12V} plasmids were used.⁽⁷⁾ For chronic liver inflammation, mice received a diet containing 0.1% DDC (Bio-Serv, Flemington, NJ) or were treated intraperitoneally twice a week with CCl₄ (Sigma) diluted 1:10 in mineral oil (Sigma) at 5 μ L/g body weight. Mice were kept in accordance with the animal regulations at the National Cancer Institute/National Institutes of Health (Bethesda, MD).

ARTICLE INFORMATION:

From the ¹Laboratory of Experimental Carcinogenesis, Center for Cancer Research, National Cancer Institute, National Institutes of Health, Bethesda, MD; ²Institute of Pathology, University Hospital of Basel, Basel, Switzerland; ³Department of Medicine I, Johannes Gutenberg University, Mainz, Germany; ⁴Biotech Research and Innovation Centre, University of Copenhagen, Copenhagen, Denmark; ⁵Cancer and Inflammation Program, National Cancer Institute-Frederick, Frederick, MD; ⁶National Cancer Institute, CCR at Leidos Inc., National Institutes of Health, Bethesda, MD; ⁷Oral and Pharyngeal Cancer Branch, National Institute of Dental and Craniofacial Research, National Institutes of Health, Bethesda, MD; ⁸Institute of Surgical Pathology, University Hospital Zurich, Zurich, Switzerland; ⁹Institute of Virology, Technische Universität München/Helmholtz Zentrum München, Munich, Germany; ¹⁰Division of Chronic Inflammation and Cancer, German Cancer Research Center, Heidelberg, Germany

ADDRESS CORRESPONDENCE AND REPRINT REQUESTS TO:

Matthias S. Matter
Institute of Pathology
University Hospital of Basel
CH-4003 Basel, Switzerland

E-mail: matthias.matter@usb.ch
Tel: +41 61 328 64 71;
Fax: +41 61 265 31 94.

STATISTICS

Data were analyzed using the Student *t* test. *P* values <0.05 were considered significant.

MICROARRAY DATA ANALYSIS

Microarray analysis was performed from four mice per tumoral group and three mice per nontumoral and control group. RNA was isolated from fresh frozen tissue using the RNeasy kit from Qiagen (Valencia, CA) and pooled from three to four histologically confirmed hepatocellular tumors per mouse and one to two locations of nontumoral tissue and control liver tissue of age-matched and sex-matched mice on normal chow. RNA was hybridized on Sentrix Mouse Ref-8v2.0 expression BeadChips (Illumina, San Diego, CA) and scanned on an iSCAN (Illumina). Data preprocessing and quantile normalization with background subtraction were performed with GenomeStudio2010 (Illumina). A signal intensity with a detection *P* > 0.05 was treated as a missing value, and only genes represented in at least 50% of all samples were included. Data analysis was performed with Qlucore (Lund, Sweden). Functional analyses were performed by gene set enrichment analysis (GSEA) and Ingenuity Pathway Analysis (IPA8.7; Ingenuity Systems, Redwood City CA). All microarray data were submitted to Gene Expression Omnibus (GSE77503). If not stated otherwise, significantly different expressed genes between two groups were identified using a *P* value of 0.001 and a fold-change difference of 1.5.

See the Supporting Information for additional information.

Results

DDC AND CCL₄ EXPOSURE CAUSED CHRONIC LIVER INFLAMMATION

Liver tumors were engineered by myristoylated and therefore activated AKT and activated truncated CAT, introduced into mouse hepatocytes by hydrodynamic tail vein injection.⁽⁸⁾ Three days after injection, mice were exposed to a DDC diet (AKT-CAT+DDC group) or kept on standard chow (AKT-CAT group). The DDC diet was tolerated well by all mice and caused typical liver pathology, which included progressive protoporphyrin with pigment deposition, formation of Mallory-Denk bodies, and hepatomegaly (data not

shown).⁽⁵⁾ In addition, DDC affected the portal area, where it caused a pronounced ductular reaction of biliary cells and a chronic inflammation containing CD3⁺ T cells and Ly6.G⁺ neutrophils (Fig. 1). Mice under continuous DDC treatment developed moderate fibrosis in the portal tract with few portal-to-portal septa, which started to form after 4–6 weeks of diet onset and gave a nodular appearance (Fig. 1, sirius red stain). Alternatively, chronic inflammation was induced by CCl₄ administration. Three days after hydrodynamic injection of AKT-CAT plasmids, mice received the first intraperitoneal injection of CCl₄, which was continued thereafter twice a week (AKT-CAT+CCl₄ group). Chronic exposure to CCl₄ caused centrilobular and portal inflammation, which consisted mainly of Ly6.G⁺ neutrophil granulocytes and CD3⁺ T lymphocytes (Fig. 1). Continuous CCl₄ administration provoked mild to moderate fibrosis around central veins with few central-to-central septa (Fig. 1, sirius red stain) starting after 4–6 weeks of treatment onset. No tumor nodules were found 4–6 weeks after hydrodynamic injection, and typical DDC or CCl₄ liver pathology was independent from stable AKT-CAT expression.

CHRONIC INFLAMMATION INDUCED BY DDC AND CCL₄ ENHANCED GROWTH OF AKT-CAT TUMORS

On average, 17.9 ± 3.4 weeks after hydrodynamic injection, AKT-CAT mice developed ascites due to excessive liver tumor growth or showed palpable liver nodules and had to be sacrificed. Macroscopic analysis revealed numerous liver tumor nodules in AKT-CAT mice (*n* = 8) (Fig. 2A). DDC treatment considerably accelerated tumor growth, and AKT-CAT+DDC mice (*n* = 7) had to be sacrificed at 11.0 ± 2.8 weeks after hydrodynamic injection (Fig. 2A). By this time, AKT-CAT+DDC tumors reached up to 5 mm in diameter, whereas the maximal diameter of the age-matched AKT-CAT tumors was <1 mm (Fig. 2A). Similarly, growth of AKT-CAT tumors was enhanced in the inflammatory environment generated by CCl₄. AKT-CAT+CCl₄ mice had to be sacrificed 13.6 ± 0.8 weeks (*n* = 4) after hydrodynamic injection, when liver tumor nodules were up to 4 mm in diameter (Fig. 2A).

Interestingly, in AKT-CAT+DDC mice, there were fewer tumors, but they were larger compared to those in AKT-CAT and AKT-CAT+CCl₄ mice (Fig. 2B). This suggests that DDC interfered with

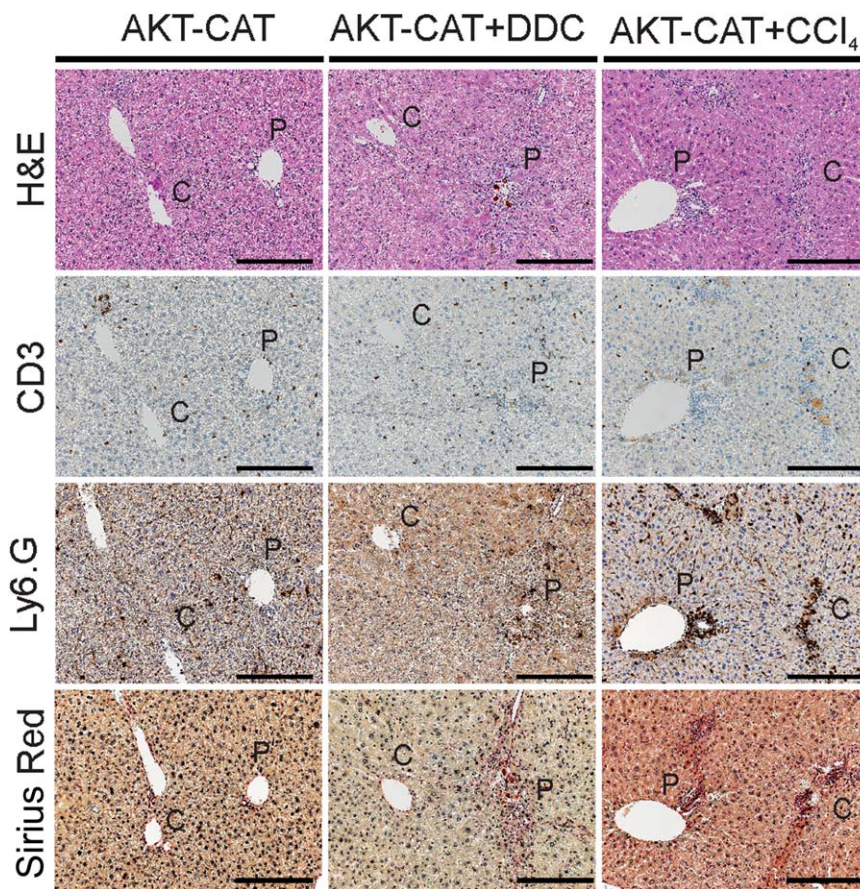


FIG. 1. Chronic liver inflammation caused by DDC and CCl_4 treatment. Three days after hydrodynamic injection of AKT-CAT plasmids, mice remained on standard chow (AKT-CAT), were subjected to a DDC diet (AKT-CAT+DDC), or were treated with intraperitoneal CCl_4 administration (AKT-CAT+ CCl_4). Hematoxylin and eosin, immunohistochemistry for CD3 and Ly6.G, and sirius red after at least 6 weeks of DDC or CCl_4 treatment. For all pictures one representative image is shown from each group (four to nine mice per group). Bar = 100 μm . Abbreviations: C, central vein; H&E, hematoxylin and eosin; P, portal tract.

tumor initiation while promoting growth of the established tumors. Of note, control mice maintained on the DDC diet ($n = 7$) for 26 weeks did not develop any neoplastic lesions. Similarly, mice treated continuously with CCl_4 ($n = 8$) for 20 weeks did not generate tumors. In addition, no tumor development was detected after injection of AKT or CAT alone to either untreated mice ($n = 6$ for each oncogene, 26 weeks of observation) or DDC-treated mice ($n = 3$ mice/group, 12 weeks of observation).

Immunohistochemistry for CAT showed strong nuclear staining with a similar intensity in all three types of AKT-CAT tumors (Fig. 2C). Likewise, the staining intensity for the second oncogene, activated AKT (phosphorylated AKT473), was also comparable in all AKT-CAT tumors, although it was considerably weaker (Fig. 2C). This confirmed a similar expression of both oncogenes in all three different AKT-CAT tumor models. To exclude that DDC diet or CCl_4 exposure changed the distribution or type of *in vivo*

transfected liver cells, we performed hydrodynamic injection with a green fluorescent protein (GFP)-positive plasmid, followed by treatment with DDC or CCl_4 for 7 and 12 days. Consistently, only hepatocytes showed GFP signals, regardless of treatment (Supporting Fig. S1). GFP-expressing hepatocytes were primarily located in the central and pericentral areas (zones 1 and 2) of the hepatic lobule. Cholangiocytes and progenitor cells did not display GFP positivity and, therefore, as shown before, were not transfected after hydrodynamic injection.⁽⁹⁾ There were no quantitative differences in GFP expression between untreated and DDC-treated mice on days 7 and 12 after hydrodynamic injection. Mice treated with CCl_4 showed a slight reduction in the number of GFP-positive hepatocytes, most likely due to the pericentral necrosis caused by the first CCl_4 injection.⁽⁶⁾ Furthermore, we compared the integrated AKT plasmid copy number in AKT-CAT, AKT-CAT+DDC, and AKT-CAT+ CCl_4 tumors. Quantitative polymerase chain

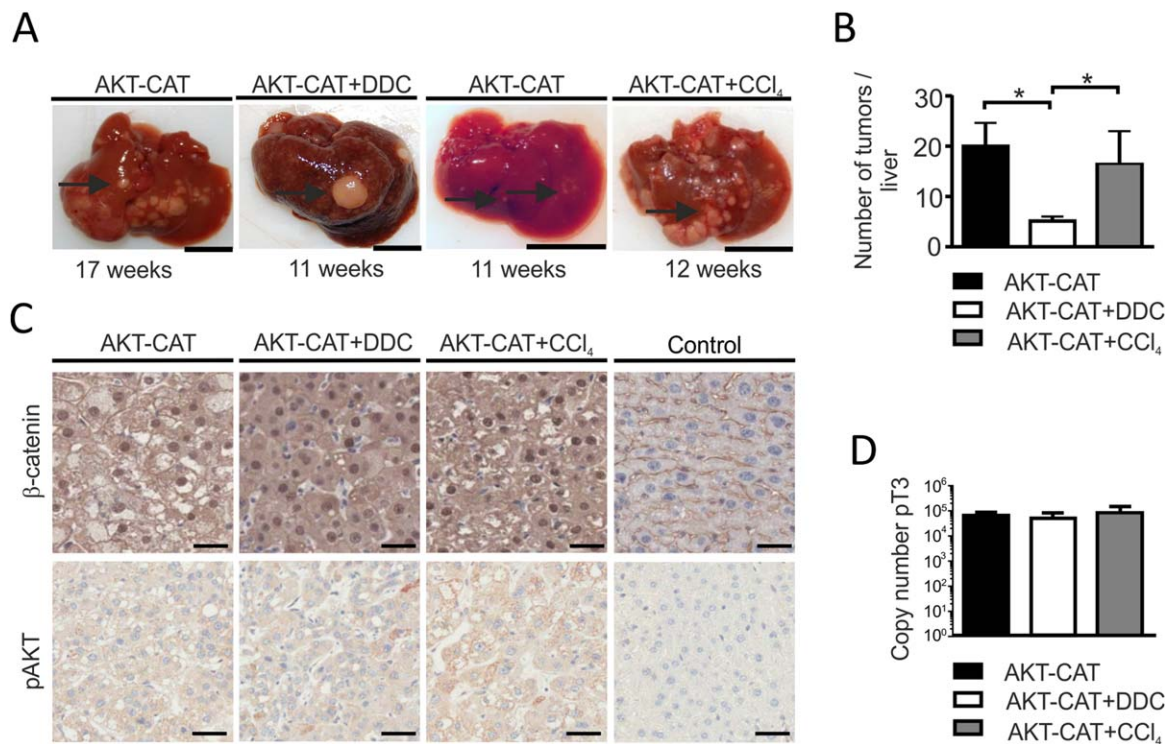


FIG. 2. AKT-CAT tumors generated in the background of chronic liver inflammation. (A) Representative macroscopic liver pictures containing AKT-CAT, AKT-CAT+DDC, or AKT-CAT+CCl₄ tumors in weeks after hydrodynamic injection. Bar = 1 cm (four to nine mice per group). Arrow indicates tumor nodule. (B) Histogram with number of liver tumor nodules with a diameter >2 mm and visible from the outside at the time of sacrifice. Mean and standard error of the mean from four to nine mice/group. (C) AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ tumors analyzed for β-catenin and p-AKT by immunohistochemistry. Bar = 50 μm. (D) Total copy number of pT3 plasmid/DNA in AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ tumors. Mean and standard error of the mean of six samples per group. For all pictures, one representative image is shown from each group.

reaction showed that integrated AKT plasmid copy number was very similar in all three tumor groups (Fig. 2D). Further, liver cell lines (SNU-449 and PLC/PRF/5) treated with DDC did not show changes in proliferation rate (Supporting Fig. S2). Altogether, these results indicate that the inflammatory microenvironment, rather than increased plasmid integration, change in cell type transfected, or direct effect of DDC on hepatocytes, was responsible for the promotion of tumor growth.

CHANGE OF TUMOR PHENOTYPE IN THE BACKGROUND OF CHRONIC LIVER INFLAMMATION

We next performed a more detailed histopathological assessment of tumor lesions according to a recent consensus report on murine liver tumor classification, which differs from human liver tumor classification.⁽¹⁰⁾

Three types of AKT-CAT driven liver tumors were recognized: hepatocellular adenoma (HCA), mixed HCA-HCC, and hepatoblastoma. A mixed HCA-HCC was diagnosed if at least 5% of the HCA was composed of an HCC. In contrast to human liver tumors, a focus of an HCC within an HCA is a common feature in murine liver tumors, thought to reflect the conversion of an HCA to a more malignant HCC phenotype.^(10,11) The predominant tumor type in all three groups was either HCA or mixed HCA-HCC (Fig. 3A). Pure HCC was not detected, and no metastasis was found in any of the tumor groups. Chronic inflammation increased the frequency of liver tumors with a mixed phenotype from 19% in untreated mice to 46% and 27% in DDC-treated and CCl₄-treated mice, respectively (Fig. 3A). Chronic inflammation also caused a marked expansion of the HCC component within the mixed HCA-HCCs. In AKT-CAT mice, the percentage of mixed HCA-HCC ranged between 5% and 25%, whereas it was 5%-50%

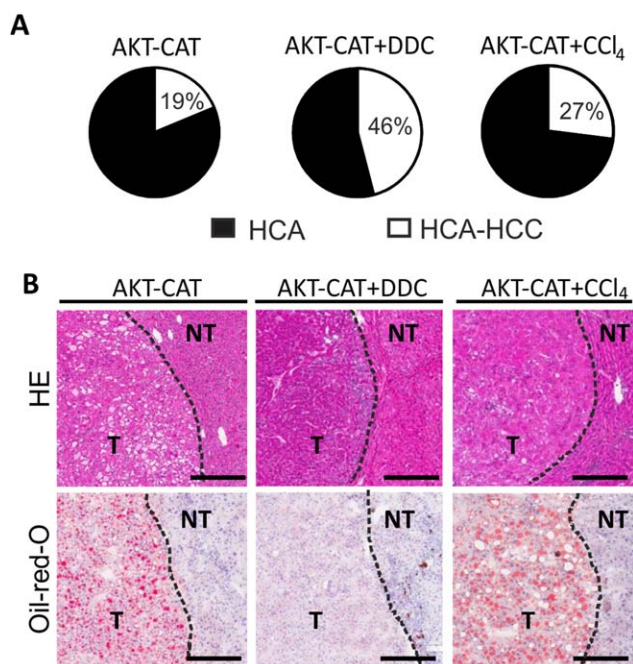


FIG. 3. Histological analysis of AKT-CAT tumors generated in the background of chronic liver inflammation. (A) Distribution of HCA and mixed HCA-HCC (with percentage) within total number of hepatocellular tumors. (B) Hematoxylin and eosin and oil red O of AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ tumors. Bar = 50 μ m. One representative picture of four to nine mice per group. Abbreviations: HE, hematoxylin and eosin; NT, nontumoral tissue; T, tumoral tissue.

and 5%-60% in AKT-CAT+DDC and AKT-CAT+CCl₄ mice, respectively. Thus, chronic inflammation promoted the progression of murine HCA into HCC. In addition, a minority of tumors was classified as hepatoblastoma. Hepatoblastoma were found only in AKT-CAT and AKT-CAT+DDC mice without any difference in morphology. Therefore, for adequate comparison between all three AKT-CAT tumor groups, further analyses were performed only on hepatocellular tumors (HCA and mixed HCA-HCC).

In accordance with earlier observations,⁽⁸⁾ most AKT-CAT tumors ($84.1 \pm 9.7\%$) were steatotic and contained lipid droplets as confirmed by oil-red O staining (Fig. 3B). Tumors were defined as steatotic if >50% of the tumor cells showed lipid droplets. In contrast, none of the AKT-CAT+DDC tumors were steatotic, but $69.8 \pm 10.3\%$ of the AKT-CAT+CCl₄ tumors were steatotic (Fig. 3B, hematoxylin and eosin and oil red O). In contrast, the number of tumor infiltrating CD3⁺ T cells was similar in all AKT-CAT

tumors, and only a few single intratumoral Ly6.G⁺ neutrophils could be found (Supporting Fig. S3).

CHANGE OF TUMOR TRANSCRIPTOME IN THE BACKGROUND OF CHRONIC LIVER INFLAMMATION

We performed transcriptome analysis of the AKT-CAT tumors developed with an inflammatory microenvironment modified by either the DDC diet or CCl₄ exposure. All tumors selected for microarray analysis had a similar size (3.0-3.4 mm in diameter) and were examined by histology to ensure that only hepatocellular tumors were included. Analysis was performed with four mice per experimental group, and RNA was pooled after isolation from three to four tumor nodules from each mouse. Unsupervised hierarchical clustering (average linkage) of all detected genes ($n = 9037$) clearly subdivided tumoral and nontumoral tissue into distinct clusters according to the experimental groups (Fig. 4A; Supporting Fig. S4). This confirmed that background inflammation induced drastic changes of the tumor transcriptome. Next, we identified genes differentially expressed in each tumor group in comparison to normal untreated liver. In total, 336 genes were differentially expressed in AKT-CAT tumors, 630 genes in AKT-CAT+DDC tumors, and 572 genes in AKT-CAT+CCl₄ tumors (Supporting Tables S1-S3). To characterize the pathways most affected by DDC and CCl₄ treatment, we used Ingenuity Pathway Analysis (IPA). In each group we identified the top five pathways associated with disease and function and searched for the differences between the tumor groups. "Cell death and survival" and "cancer" were shared among all AKT-CAT tumors (Table 1); however, *P* values were higher in DDC-treated and CCl₄-treated tumors. "Cellular growth and proliferation" exemplified the AKT-CAT+DDC and AKT-CAT+CCl₄ tumors, while in AKT-CAT tumors this function was ranked only seventeenth. Accordingly, cell proliferation as measured by Ki-67 confirmed that tumor growth rate was considerably higher in AKT-CAT+DDC and AKT-CAT+CCl₄ tumors compared to AKT-CAT tumors (Fig. 4B). The remaining most affected pathways were unique for each group (see Table 1). Thus, the background inflammation had a strong impact on pathway activation in each AKT-CAT tumor group.

Next, we analyzed the "cellular growth and proliferation" network, which contained 205 genes for

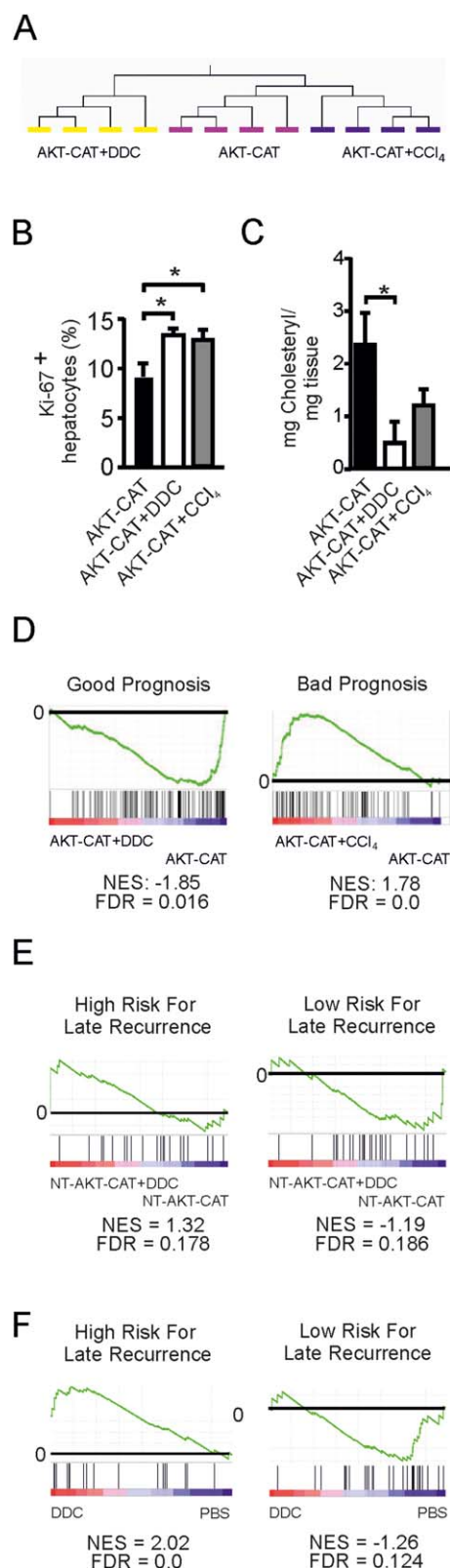


FIG. 4

AKT-CAT+DDC tumors and 169 genes for AKT-CAT+CCl₄ tumors (Supporting Tables S4 and S5). Comparison between these two gene sets identified Col4a2, Cxcl16, Elf3, Galectin-3, Lxn, Myp, and Nedd9, which were shared by AKT-CAT+DDC and AKT-CAT+CCl₄ tumors and which showed furthermore a significant deregulation from AKT-CAT tumors ($P < 0.01$, fold change > 1.5). As Cxcl16, Galectin-3, and Nedd9 have been associated with worse prognosis in HCC patients,⁽¹²⁻¹⁴⁾ we tested their expression by quantitative polymerase chain reaction (Supporting Fig. S5). Indeed, their expression was highest in AKT-CAT+DDC and AKT-CAT+CCl₄ tumors.

AKT-CAT+DDC tumors also showed an increase in pathways connected to vascular disease, which included atherosclerosis and arteriosclerosis. AKT-CAT+CCl₄ tumors showed a significant change in cardiovascular system development and function, which included proliferation and migration of endothelial cells as well as neovascularization. Yet, microvessel density remained similar in all AKT-CAT tumor groups (Supporting Fig. S6).

Histologically, one of the most striking characteristics of DDC-treated tumors was loss of lipid droplets. IPA showed that lipid metabolism was more significantly changed in AKT-CAT+DDC tumors ($P = 5.62E08$ - $2.46E-3$) than in AKT-CAT tumors ($P = 1.71E03$ - $1.33E02$) and AKT-CAT+CCl₄ tumors ($P = 5.53E04$ - $9.67E03$). Detailed analysis showed that synthesis, flux, efflux, and uptake of lipids, as well as uptake and metabolism of fatty acid, were affected in AKT-CAT+DDC tumors and that the majority of these functions showed a reduced activation score. To corroborate the involvement in lipid metabolism further, we determined the content of cholesteryl esters

FIG. 4. Transcriptome of AKT-CAT tumors generated in the background of chronic liver inflammation. (A) Unsupervised hierarchical clustering of gene expression profiles from AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ tumors. (B) Percentage of proliferating Ki-67-positive hepatocytes within total hepatocytes, measured by immunohistochemistry. Mean and standard error of the mean from seven to eight tumors per group. (C) Cholesteryl ester concentration in AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ tumors. Mean and standard error of the mean of four tumor samples per group. (D) GSEA of AKT-CAT and AKT-CAT+DDC tumors using a gene set with good prognosis (left) and of AKT-CAT and AKT-CAT+CCl₄ tumors using a gene set with bad prognosis (right). (E,F) GSEA of non-tumoral tissue of AKT-CAT and AKT-CAT+DDC mice (E) or mice treated with DDC only (F) using gene sets with low and high risks of late recurrence. Abbreviations: FDR, false discovery rate; NES, normalized enrichment score.

TABLE 1. Most Significant Diseases and Function Affected in AKT-CAT, AKT-CAT+DDC, and AKT-CAT+CCl₄ Tumors

Tumor group	<i>P</i>
<i>Shared by all tumor groups</i>	
AKT-CAT	
Cell death and survival	6.68E-07-1.33E-02
Cancer	2.32E-06-1.34E-02
AKT-CAT+DDC	
Cell death and survival	5.2E-14-2.29E-03
Cancer	6.03E-11-2.46E-03
AKT-CAT+CCl ₄	
Cell death and survival	1.82E-09-1.04E-02
Cancer	2.33E-06-1.05E-02
<i>Shared by AKT-CAT+DDC and AKT-CAT+CCl₄</i>	
AKT-CAT+DDC	
Cellular growth and proliferation	1.75E-13-2.23E-03
AKT-CAT+CCl ₄	
Cellular growth and proliferation	5.12E-10-1.06E-02
<i>Unique in each tumor group</i>	
AKT-CAT	
Cell signaling	6.7E-07-3.53E-03
Tumor morphology	2.32E-06-1.33E-02
Organismal injury and abnormalities	1.28E-05-1.33E-02
AKT-CAT+DDC	
Cardiovascular disease	2.01E-10-2.29E-03
Development disorder	1.17E-09-4.97E-04
AKT-CAT+CCl ₄	
Cardiovascular system development and function	6.19E-08-1.02E-02
Organismal development	6.19E-08-1.02E-02

and triglycerides, which are both stored in lipid droplets.⁽¹⁵⁾ Cholesteryl ester concentration was significantly reduced in AKT-CAT+DDC tumors in comparison to AKT-CAT tumors, whereas AKT-CAT+CCl₄ tumors had an intermediate concentration (Fig. 4C). Triglycerides were only slightly reduced in AKT-CAT+DDC tumors (Supporting Fig. S7). Accordingly, proteins which are involved in fatty acid and cholesterol metabolism⁽¹⁶⁾—fatty acid synthase, phospho-adenosine triphosphate-citrate lyase, mammalian long chain acyl-coenzyme A synthetase, cytoplasmic acetyl-coenzyme A synthetase, and phospho-S6-ribosomal protein—showed reduced expression in AKT-CAT+DDC and AKT-CAT+CCl₄ tumors (Supporting Fig. S7).

GSEA OF TUMORAL AND NONTUMORAL TISSUE

Next, we performed cross-species comparison and GSEA to relate our murine data to human data.⁽¹⁷⁾ We used our previously published human HCC gene sets, which define HCC patients with good and bad prognosis.⁽¹⁸⁾ Consistent with a more aggressive phenotype of tumors grown in the inflammatory environment, AKT-CAT+DDC tumors were characterized

by a significant decrease of good prognosis genes (Fig. 4D) and AKT-CAT+CCl₄ tumors showed an enrichment of poor prognosis genes (Fig. 4D). Thus, the biology of AKT-CAT+DDC and AKT-CAT+CCl₄ tumors showed similarities with human HCCs.

Given the importance of the nontumoral liver tissue in HCC prognosis, we performed transcriptome analysis of the nontumoral murine liver tissue by microarray and executed GSEA with gene sets from HCC patients with low or high risk of HCC recurrence.⁽¹⁹⁾ Nontumoral liver tissue from AKT-CAT+DDC mice showed enrichment of genes with high risk of late recurrence and, vice versa, the genes with low risk of late recurrence were enriched in AKT-CAT nontumoral liver tissue (Fig. 4E). Therefore, the worse outcome of tumor-bearing mice under DDC treatment was deducible not only from tumoral transcriptome but also from the nontumoral transcriptome, similar to HCC patients. Interestingly, liver tissue from tumor-free mice treated only with DDC revealed an even further enrichment of genes related to worse prognosis (Fig. 4F). Therefore, AKT-CAT tumors provoked changes in the nontumoral liver tissue, which improved prognosis. In contrast, there was no predictive value of the nontumoral liver tissue from CCl₄-treated mice using these gene sets at a false discovery rate of <0.25.

IMPACT OF DDC TREATMENT ON AKT-NRAS^{G12V}-DRIVEN HEPATOCARCINOGENESIS

Furthermore, we asked whether a tumor driven by a different combination of oncogenes would be similarly affected by chronic liver inflammation. To address this issue, we generated liver tumors by hydrodynamic injection of AKT and mutated NRAS^{G12V} and placed the mice on either a DDC diet (AKT-NRAS^{G12V}+DDC) or normal chow (AKT-NRAS^{G12V}). AKT-NRAS^{G12V} tumors were more aggressive than AKT-CAT tumors, and AKT-NRAS^{G12V} mice had to be sacrificed 10 weeks after hydrodynamic injection (versus 17.9 weeks for AKT-CAT model). AKT-NRAS^{G12V}+DDC mice had to be sacrificed slightly earlier, at 8.4 weeks after hydrodynamic injection, mainly due to ascites formation. Most of the liver tissue was occupied by tumor nodules in both groups at the time of sacrifice (Fig. 5A). DDC treatment did not significantly influence either the number of AKT-NRAS^{G12V} tumor nodules (Fig. 5B) or the proliferation rate as measured by immunohistochemistry

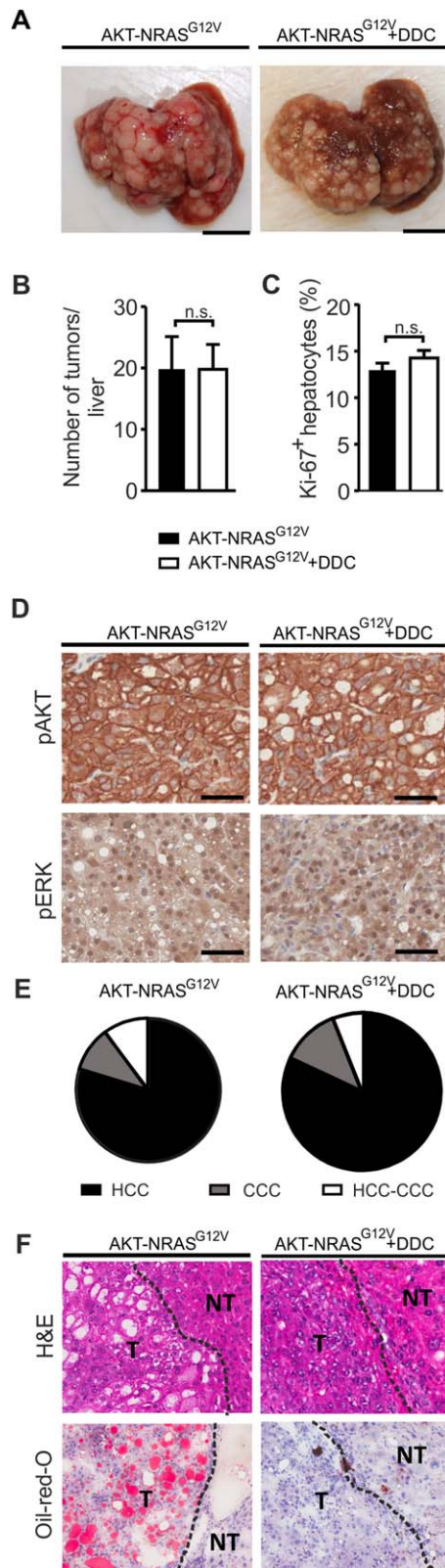


FIG. 5

for Ki-67 (Fig. 5C). Oncogene expression was similar in both tumor groups, as measured by the expression of extracellular signal-regulated kinase, a downstream target of RAS activation and phosphorylated AKT (Fig. 5D). Histologically, the majority of AKT-NRAS^{G12V} tumors were classified as HCC.⁽⁷⁾ As shown,⁽⁷⁾ we also found cholangiocarcinomas (CCCs) or mixed HCC-CCC (Supporting Fig. S8). The DDC diet did not change the relative proportion of HCC, CCC, and mixed HCC-CCC, which varied 80%-82%, 10%-12%, and 6%-10% in the two groups (Fig. 5E). However, analogous to AKT-CAT tumors, DDC treatment caused a reduction in lipid accumulation in (Fig. 5F), and average percentage of steatotic HCCs decreased from 56.1% in AKT-NRAS^{G12V} HCCs to 13.5% in AKT-NRAS^{G12V}+DDC tumors (data not shown). No metastasis was found in any of the tumor groups.

Transcriptome of histologically confirmed HCCs with similar size from AKT-NRAS^{G12V} and AKT-NRAS^{G12V}+DDC mice was analyzed. Unsupervised hierarchical clustering of all detected genes ($n = 9037$) clearly separated tumors into two clusters determined by DDC treatment (Supporting Fig. S9). In total, 1629 genes were differentially expressed in AKT-NRAS^{G12V} tumors and 1913 genes in AKT-NRAS^{G12V}+DDC tumors in comparison to normal untreated liver tissue. IPA of the differentially expressed genes showed that the pathways affected were mostly shared in both experimental groups (Table 2). However, P values were always higher in AKT-NRAS^{G12V}+DDC tumors, indicating that DDC mainly enhanced the differences of the changed pathways. Yet, cross-species comparison and GSEA with our published human gene sets⁽¹⁸⁾ further demonstrated an enrichment of poor prognosis gene sets in AKT-NRAS^{G12V}+DDC tumors and of

FIG. 5. (A) Macroscopic liver pictures with AKT-NRAS^{G12V} or AKT-NRAS^{G12V}+DDC tumors. Bar = 1 cm. (B) Number of liver tumor nodules >2 mm in diameter visible from outside in AKT-NRAS^{G12V} or AKT-NRAS^{G12V}+DDC mice at the time of sacrifice. Mean and standard error of the mean from three to four mice per group. (C) Percentage of Ki-67-positive hepatocytes within total hepatocytes, measured by immunohistochemistry. Mean and standard error of the mean of six to nine tumor samples/group. (D) Immunohistochemistry for phosphorylated AKT and phosphorylated extracellular signal-regulated kinase in AKT-NRAS^{G12V} and AKT-NRAS^{G12V}+DDC tumors. (E) Distribution of HCC, CCC, and mixed HCC-CCC in AKT-NRAS^{G12V} and AKT-NRAS^{G12V}+DDC tumors from three to four mice per group. (F) Hematoxylin and eosin and oil red O staining of AKT-NRAS^{G12V} and AKT-NRAS^{G12V}+DDC tumors. Abbreviations: H&E, hematoxylin and eosin; n.s., nonsignificant; NT, nontumoral tissue; pERK, phosphorylated extracellular signal-regulated kinase; T, tumoral tissue.

TABLE 2. Most Significant Diseases and Function Affected in AKT-NRAS^{G12V} and AKT-NRAS^{G12V}+DDC Tumors

Tumor group	<i>P</i>
<i>Shared pathways</i>	
AKT-NRAS ^{G12V}	
Cellular growth and proliferation	4.73E-22-4.11E-04
Cell death and survival	1.51E-16-6.42E04
Cardiovascular system development and function	2.17E-15-6.42E-04
Cancer	3.19E-14-6.16E04
AKT-NRAS ^{G12V} +DDC	
Cellular growth and proliferation	4.17E-30-4.02E-05
Cell death and survival	8.54E-29-5.3E-05
Cancer	4.66E-21-5.14E-05
Cardiovascular system development and function	1.44E-18-4.21E-05
<i>Unique pathways</i>	
AKT-NRAS ^{G12V}	
Organismal survival	5.95E-17-1.83E-04
AKT-NRAS ^{G12V} +DDC	
Organismal injury and abnormalities	4.66E-21-5.2E-05

good prognosis gene sets in AKT-NRAS^{G12V} tumors (Supporting Fig. S9), further supporting that biology of AKT-NRAS^{G12V}+DDC tumors reflected human HCC.

Finally, we wanted to compare effects exerted by DDC on AKT-CAT tumors with the DDC effects on AKT-NRAS^{G12V} tumors. A bioequivalence test⁽²⁰⁾ of similarities at a fold change >1.5 and *P* < 0.05 revealed that 90.1% of the genes were similarly expressed between AKT-CAT and AKT-CAT+DDC tumors and 94.9% between AKT-NRAS and AKT-NRAS^{G12V}+DDC tumors. Therefore, DDC had a more pronounced effect on AKT-CAT tumors than on AKT-NRAS^{G12V} tumors. For direct comparison of DDC effects we normalized AKT-CAT+DDC to AKT-CAT and AKT-NRAS^{G12V}+DDC to AKT-NRAS^{G12V} and compared the two groups. We identified a total of 393 genes significantly changed between normalized AKT-CAT+DDC and AKT-NRAS^{G12V}+DDC data sets. IPA showed that the most significantly changed pathway by DDC in AKT-CAT tumors in comparison to AKT-NRAS^{G12V} tumors was “cellular growth and proliferation” (Supporting Table S6), followed by “cellular assembly and organization,” “cardiovascular system development and function,” “organismal survival,” and “cancer.” Therefore, DDC had a different effect on AKT-CAT tumors than on AKT-NRAS^{G12V} tumors.

Discussion

Several studies have revealed the importance of the inflammatory liver microenvironment in HCC development. For example, it has been shown that prognosis of HCC patients can be predicted by the gene

signature of nontumoral liver tissue.^(19,21) Furthermore, HCC patients with an increase of T helper 2 cytokines in relation to T helper 1 cytokines in nontumoral liver tissue revealed increased venous metastasis and poor prognosis.⁽²²⁾ However, many of the mechanisms linking inflammation to tumor development remain unclear, and little is known as to what extent chronic liver inflammation contributes to HCC phenotype. In this study, we developed an experimental mouse model, in which murine liver tumors were driven by specific oncogenes and developed in different inflammatory conditions. Chronic inflammation promoted tumor growth and aggressiveness but had a stronger effect on AKT-CAT tumors than on AKT-NRAS^{G12V} tumors. Therefore, liver tumor phenotype is dependent on the interplay between oncogenic driver genes and the microenvironment.

Our study goes along with earlier murine studies, which also described a tumor promoting effect by DDC or CCl₄.^(23,24) We noted that both DDC and CCl₄ affected especially “cell growth and proliferation” in AKT-CAT tumors and that Galectin-3, Nedd9, and Cxcl16 were among several genes specifically up-regulated. These three genes have been associated additionally with worse prognosis in human HCC.⁽¹²⁻¹⁴⁾ Galectin-3 has been shown to promote tumor progression and metastasis formation in HCC.⁽²⁵⁾ Interestingly, Galectin-3 has also been reported to be up-regulated in injured liver⁽²⁶⁾ and may therefore link chronic liver damage and tumor development. Similarly, Cxcl16 and its receptor CXCR16, as well as Nedd9, have been shown to be involved in HCC progression and metastasis formation.^(27,28) AKT-CAT+DDC and AKT-CAT+CCl₄ tumors also showed a higher proportion of HCCs, and GSEA confirmed that changes in tumoral and nontumoral tissue reflected human HCC biology. Interestingly, DDC treatment induced loss of lipid droplets with a reduction in cholesteryl content in AKT-CAT tumors. Accordingly, several genes involved in fatty acid and cholesterol metabolism were reduced in AKT-CAT+DDC and AKT-CAT+CCl₄ tumors in comparison to AKT-CAT tumors.

Several effects appreciated in AKT-CAT tumors under DDC treatment, such as lipid loss and a decreased expression of genes associated with good survival by GSEA, were also observed in AKT-NRAS^{G12V}+DDC tumors. However, in line with the bioequivalence test, the impact of DDC treatment on AKT-NRAS^{G12V} tumors was less pronounced and was primarily an enhancement of changes already procured by AKT-NRAS^{G12V}. For example, the effect on

pathways involved in cell growth and proliferation was negligible. Direct comparison between AKT-CAT and AKT-NRAS^{G12V} tumors also revealed that DDC had a different effect on AKT-CAT tumors compared to AKT-NRAS^{G12V} tumors. As expected, the effect on “cell growth and proliferation” was the most significant difference.

The differences in the response to inflammatory stimuli between AKT-CAT and AKT-NRAS^{G12V} tumors may be explained in two ways. First, untreated AKT-NRAS^{G12V} tumors were more proliferative, grew significantly faster, and had a more aggressive phenotype than untreated AKT-CAT tumors. This suggests that fast-growing malignant tumors are less susceptible to the inflammatory microenvironment and more likely to survive under adverse conditions. Second, there may be oncogene-specific differences in the response to cytokines or chemokines produced by the tumor microenvironment. Thus, it has been recently shown that CAT has an intrinsic proinflammatory transcriptional program.⁽²⁹⁾ Therefore, AKT-CAT tumors may be more susceptible to inflammatory stimuli than AKT-NRAS^{G12V} tumors.

Although we demonstrated that chronic inflammation plays an important role in tumor phenotype, our results do not allow differentiation of whether the phenotype is defined by the best-fit tumor cells capable of surviving in the background inflammation or by the plasticity of tumor cells with transformation into a more aggressive phenotype under the pressure of chronic inflammation. DDC-treated mice showed a reduction in the number of AKT-CAT tumor nodules, suggesting that only selective tumor cells had the ability to survive in the background of chronic inflammation driven by DDC and that some tumor cells may have been eliminated by chronic inflammation. This notion is supported by the fact that chronic inflammation can either promote or inhibit tumor growth.⁽²⁾ In contrast, the number of AKT-CAT tumors was similar in untreated and CCl₄-treated mice, indicating that inflammation caused by CCl₄ promoted a transformation into a more aggressive phenotype. Therefore, the appreciated change in phenotype of AKT-CAT tumors was most likely a combination of both selection and transformation of tumor cells.⁽³⁰⁾

In conclusion, our study provides evidence that the inflammatory milieu changes the behavior and phenotype of HCC in a manner dependent on the driving oncogenes. Furthermore, tumors with the same mutations and driving oncogenes may have substantially different transcriptome depending on the type of

inflammation. As a consequence, response to systemic cancer treatment may also depend on the tumor microenvironment.

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Author names in bold designate shared co-first authorship.

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